TOPCARE ANTI DIARRHEAL- loperamide hydrochloride tablet, film coated
Topco Associates LLC

Topco Associates LLC. Anti-Diarrheal Tablets Drug Facts

Active ingredient (in each caplet)

Loperamide HCl 2 mg

Purpose

Anti-diarrheal

Use

controls symptoms of diarrhea, including Travelers' Diarrhea

Warnings

Allergy alert: Do not use if you have ever had a rash or other allergic reaction to loperamide HCl

Heart alert: Taking more than directed can cause serious heart problems or death

Do not use

if you have bloody or black stool

Ask a doctor before use if you have

- fever
- mucus in the stool
- a history of liver disease
- a history of abnormal heart rhythm

Ask a doctor or pharmacist before use if you are

taking a prescription drug. Loperamide may interact with certain prescription drugs.

When using this product

tiredness, drowsiness or dizziness may occur. Be careful when driving or operating machinery.

Stop use and ask a doctor if

- symptoms get worse
- diarrhea lasts for more than 2 days
- you get abdominal swelling or bulging. These may be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Directions

- **drink plenty of clear fluids to help prevent dehydration caused by diarrhea**
- find right dose on chart. If possible, use weight to dose; otherwise, use age.

adults and children 12 years and over	2 caplets after the first loose stool; 1 caplet after each subsequent loose stool; but no more than 4 caplets in 24 hours
children 9-11 years (60-95 lbs)	1 caplet after the first loose stool; 1/2 caplet after each subsequent loose stool; but no more than 3 caplets in 24 hours
children 6-8 years (48-59 lbs)	1 caplet after the first loose stool; 1/2 caplet after each subsequent loose stool; but no more than 2 caplets in 24 hours
children 2-5 years (34 to 47 lbs)	ask a doctor
children under 2 years (up to 33 lbs)	do not use

Other information

- store at 20°-25°C (68°-77°F)
- see end panel for lot number and expiration date

Inactive ingredients

anhydrous lactose, carnauba wax, D&C yellow no. 10 aluminum lake, FD&C blue no. 1 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, pregelatinized starch

Questions or comments?

1-888-423-0139

Principal Display Panel

COMPARE TO IMODIUM® A-D ACTIVE INGREDIENT

Anti-Diarrheal Tablets

LOPERAMIDE HYDROCHLORIDE TABLETS, 2 mg

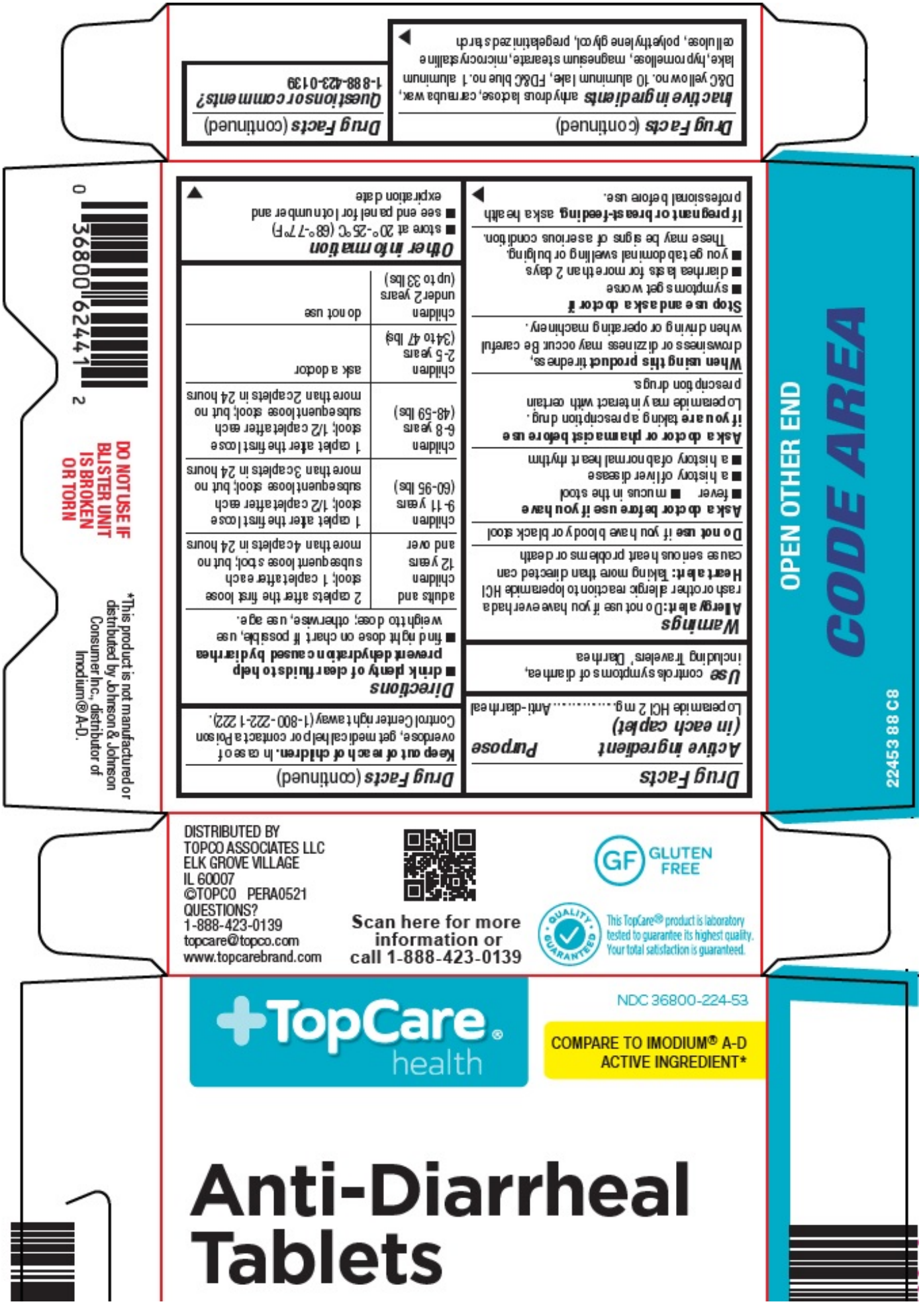
Anti-Diarrheal

● Controls the Symptoms of Diarrhea

12 CAPLETS†

†CAPSULE-SHAPED TABLETS

actual size



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Drug Facts (continued)

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OPEN OTHER END

CODE AREA

22453 88 C8

0 36800 62441 2

DO NOT USE IF BLISTER UNIT IS BROKEN OR TORN

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Imodium® A-D.

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IL 60007
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QUESTIONS?
1-888-423-0139
topcare@topco.com
www.topcarebrand.com

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GLUTEN FREE

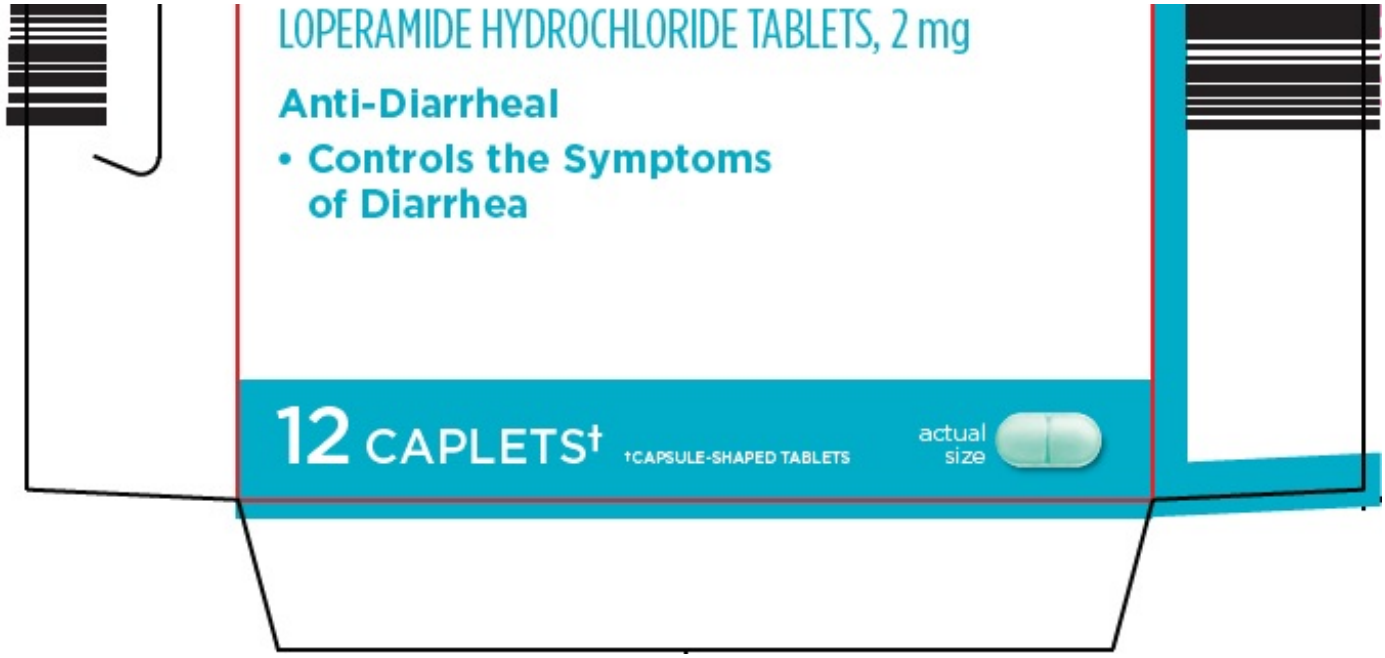
This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

TopCare® health

NDC 36800-224-53

COMPARE TO IMODIUM® A-D ACTIVE INGREDIENT*

Anti-Diarrheal Tablets



TOPCARE ANTI DIARRHEAL

loperamide hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:36800-224
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LOPERAMIDE HYDROCHLORIDE (UNII: 77TI35393C) (LOPERAMIDE - UNII:6X9OC3H4II)	LOPERAMIDE HYDROCHLORIDE	2 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

Color	GREEN	**Score**	2 pieces
Shape	OVAL	**Size**	10mm

Flavor		**Imprint Code**	L2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-224-91	6 in 1 CARTON	08/07/2003	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:36800-224-53	12 in 1 CARTON	02/26/2003	
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:36800-224-62	24 in 1 CARTON	04/22/2003	
3		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:36800-224-80	1 in 1 CARTON	11/12/2007	03/31/2020
4		96 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075232	02/26/2003	

Labeler - Topco Associates LLC (006935977)

Revised: 11/2021 Topco Associates LLC